

Isozymes and Canonical Discriminant Analysis to Identify Pistachio (*Pistacia vera* L.) Germplasm

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Abstract. Canonical discriminant analysis (CDA) of morphometric data of buds, leaves, and fruit, as well as isozyme analysis (esterase, peroxidase, and acid phosphatase) of leaf samples, were used to identify eight male pistachio selections and 10 female pistachio cultivars. According to the CDA, 77% and 93% of the total variance was summarized by the first three canonical discriminant functions for the female and male selections, respectively. Fruit characteristics, particularly fruit fresh and dry weights and fruit length, accounted for most of the discriminatory power for the female cultivars, while the dimensions of the leaves, principally leaf rachis length, were the most effective discriminating characters for the males. Isozyme analysis showed a higher degree of polymorphism in the male than the female genetic pool. Hence, using only three enzymes, it was possible to identify all of the male selections, but only 50% of the females. Peroxidase polymorphism clearly demonstrated the greater phylogenetic distance between 'Kerman' and the local cultivars, as well as between 'Cerasola', a quite different cultivar with a reddish hull, and the others tested. The combination of CDA and isozyme analysis enhanced the possibility of uniquely identifying the female cultivars.

The pistachio tree, native to western Asia and Asia Minor, was introduced into Italy from Syria in A.D. 30 by the Romans (Minà Palumbo, 1882). Its cultivation spread to Sicily during the Arab domination (A.D. 827 to 1040). Despite the long cultivation period, only a few cultivars of pistachio are grown in Sicily, where the Italian industry is located. 'Bianca' is the main cultivar, with others representing <3% of the cultivated area (Barone et al., 1985). Most of these minor cultivars are no longer grown commercially nor are they available in nurseries. Therefore, the genetic reserve is being rapidly eroded. Nevertheless, the Sicilian germplasm has some desirable characteristics, such as color, flavor, and nut quality (Caruso et al., 1986, 1987). Indeed, Sicilian pistachios have long been appreciated for the intense green of their kernels and rich, oily nutlike flavor (Woodroof, 1967).

As a part of a breeding research program carried out at the Univ. of Palermo, a germplasm collection was established in 1984, including 'Bianca' and 'Kerman' as standard cultivars, together with eight indigenous cultivars and eight unnamed male pistachio selections. As the possibility of synonymy in such a collection could not be excluded, the varietal identi-

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in Sicily, found some cases of homonymy within those accessions.

A promising method for cultivar identification and classification is offered by the numerical taxonomic approach (Rhodes et al., 1971). Some techniques of multivariate analyses have been applied widely to discriminate and classify cultivars and clones of several fruit tree species (Alessandri and Casini, 1992; Cruz-Castillo et al., 1992; Gogorcena and Ortiz, 1993; Hillig and Iezzoni, 1988; Perez-Gonzales, 1992), but no report for pistachio was found.

The objectives of this study were to assess the usefulness of a multivariate statistical technique, i.e., canonical discriminant analysis (CDA), and isozyme polymorphism for identifying pistachio cultivars and to determine if synonymy exists in our collection. We also wanted to detect the degree of similarity or dissimilarity within the Italian pistachio germplasm.

Materials and Methods

Plant material. Two well-known female cultivars, 'Bianca' (A) and 'Kerman' (J), together with eight other putative distinct female cultivars, i.e., 'Cappuccia' (B), 'Cerasola' (C), 'Ghiandalora' (D), 'Gialla' (E), 'Insolia' (F), 'Pignatone' (G), 'Silvana' (H), and 'Tardiva' (I), and eight unnamed male pistachio selections, grown in a germplasm collection at Campobello di Licata (Italy 37°30'N), were examined. The collection was established in 1984 in a randomized block design with 15 trees per cultivar and three trees per male selection. All trees were budded on *P. terebinthus* seedling rootstocks and trained to a vase shape. The main morphological and phenological characteristics of this germplasm have been described by Caruso et al. (1995).

Isozymes. Fully expanded leaves from the median part of nonbearing shoots of 9-year-old trees of each female and male selection were sampled in July and stored at -20°C up to 1 week. A 0.5-g sample of freeze-dried leaves

of this plant material was recognized as a problem of primary importance.

In the last decade, isozyme techniques have been used to identify cultivars (Torres, 1983). However, few attempts with pistachio are reported (Arulsekaran and Parfitt, 1986). Loukas and Pontikis (1979) used pollen isozymes to detect the degree of phylogenetic relationship among various species and types of *Pistacia*, but, because they were working with pollen, they were limited to studying male genotypes. More recently, Dollo (1991), using leaf samples from 22 species and cultivars of *Pistacia* grown

Table 1. Standardized canonical coefficient (SCC) and correlation coefficient (*r*) between the first three canonical discriminant functions (CDF₁, CDF₂, CDF₃) and variables of the morphometric characters of the female cultivars.

Character	Canonical discriminant function					
	CDF ₁		CDF ₂		CDF ₃	
	SCC	<i>r</i>	SCC	<i>r</i>	SCC	<i>r</i>
Leaf rachis length	-0.057	-0.001	0.239	0.259	-0.479	-0.316
Basal leaflet length	0.361	0.225	0.090	0.346	0.360	0.180
Basal leaflet width	-0.069	0.199	0.437	0.450	0.061	-0.048
Central leaflet length	-0.313	0.142	-0.440	0.216	0.431	0.119
Central leaflet width	0.085	0.182	0.159	0.444	0.155	-0.104
Apical leaflet length	0.248	0.228	-0.182	0.189	0.035	0.065
Apical leaflet width	-0.050	0.121	0.397	0.415	-0.866	-0.277
Inflorescence bud length	0.315	0.414	0.193	0.233	0.359	0.413
Inflorescence bud max diameter	0.020	0.311	-0.061	0.122	0.343	0.389
Inflorescence bud min diameter	0.157	0.340	0.259	0.213	-0.225	0.179
Inflorescence bud fresh weight	0.037	0.367	-0.061	0.194	-0.112	0.282
Inflorescence bud dry weight	0.331	0.379	0.212	0.188	0.237	0.297
Fruit length	-0.446	-0.048	0.756	0.397	0.178	0.106
Fruit maximum diameter	0.312	0.473	-0.031	-0.138	-0.275	-0.322
Fruit minimum diameter	0.422	0.503	0.260	0.054	-0.362	-0.297
Fruit fresh weight	0.737	0.421	-1.053	-0.189	-0.166	-0.106
Fruit dry weight	-0.597	0.186	0.396	-0.145	0.417	-0.010
Variance explained (%)	43.85		23.43		9.62	

was used. The extraction buffer (2 ml) contained 50 mM TRIZMA base, an antioxidant (Sigma A1179), 10 mM MgCl₂, 10 mM diethyldithiocarbamic acid, 1 mM 2-mercaptoethanol, and 1.0% polyethylene glycol (M₃₅₀₀). The pH was adjusted to 8.0 with 0.1 N HCl. Samples were homogenized in an Ultra-Turrax (Janke and Kunkel, Staufen, Germany) homogenizer for 30 sec with polyvinylpyrrolidone and centrifuged for 15 min (18,100× g) at 4C; supernatant was collected and centrifuged again. The following extract was immediately used for electrophoresis. Amount of total proteins was measured according to Bradford (1976). A total of 25 µl/sample of extract was submitted to electrophoresis in 12% polyacrylamide gel for 2.30 h at constant voltage of 120 V, 20 mA, at 4C. The gel electrophoresis apparatus used was the Mini Protean II (Bio-rad, Hercules, Calif.). The enzyme systems studied were esterase (EST), peroxidase (PER), and acid phosphatase (AcPH). Staining procedures were as described by Shaw (1970). The entire procedure was conducted three times.

The results from isozyme analysis were used for producing two matrices of dissimilarity between accessions. Based on these matrices, the grouping of the accessions was made by hierarchical cluster analyses by using normalized percent disagreement as the distance index and the complete linkage method, which uses the most distant pair of objects, to compute between-cluster distances (farthest neighbor). The hierarchy of clusters formed during cluster analysis was illustrated as a dendrogram.

Multivariate analysis. Seventeen variables for each of the 10 females and 12 variables for each of the eight male accessions were determined by at least 50 measurements for each character (Tables 1 and 2), on samples randomly collected from the entire collection. CDA was performed separately on the female and male set of the morphometric data using the MGLH procedure of the SYSTAT statistical package (Wilkinson, 1990). The discriminant power of the obtained canonical discriminant functions was tested by checking predicted vs. actual group membership of each observation, i.e., by applying the obtained canonical discriminant functions (CDFs) to the set of data and calculating the percentage of cases correctly classified. Canonical scores of major CDFs were submitted to analysis of variance (ANOVA). Means of standardized canonical scores of the major CDFs were plotted.

Results and Discussion

Isozymes. All the enzymes tested were polymorphic. Isozymes of PER were slightly polymorphic, showing three banding patterns for all female cultivars (Fig. 1) and five for male selections (Fig. 2).

Esterases. All the male and the female selections can be divided into five groups, each with identical electrophoretic patterns. Three male selections (M3, M7, and M9) and three female cultivars ('Bianca', 'Cappuccia',

Table 2. Means of standardized canonical scores of the first three canonical discriminant functions (CDF) for the female cultivars.

Cultivar	Canonical discriminant function		
	CDF ₁ ^z	CDF ₂ ^z	CDF ₃ ^z
A ^y	-0.324 c	0.515 de	-1.021 ab
B	0.795 d	0.083 cd	1.504 e
C	-0.304 c	2.524 f	-0.355 bc
D	-0.803 c	1.027 e	0.716 d
E	-0.528 c	1.021 e	-0.979 ab
F	-2.034 b	-0.283 c	-0.401 ab
G	-0.431 c	-0.102 cd	-0.341 bc
H	4.651 f	-1.889 b	-1.077 a
I	2.111 e	0.224 cd	1.693 e
J	-3.185 a	-3.187 a	0.316 cd

^zMean separation within each column by Tukey's HSD test ($P \leq 0.01$).

^yA = 'Bianca'; B = 'Cappuccia'; C = 'Cerasola'; D = 'Ghiandalora'; E = 'Giulla'; F = 'Insolia'; G = 'Pignatone'; H = 'Silvana'; I = 'Tardiva'; J = 'Kerman'.

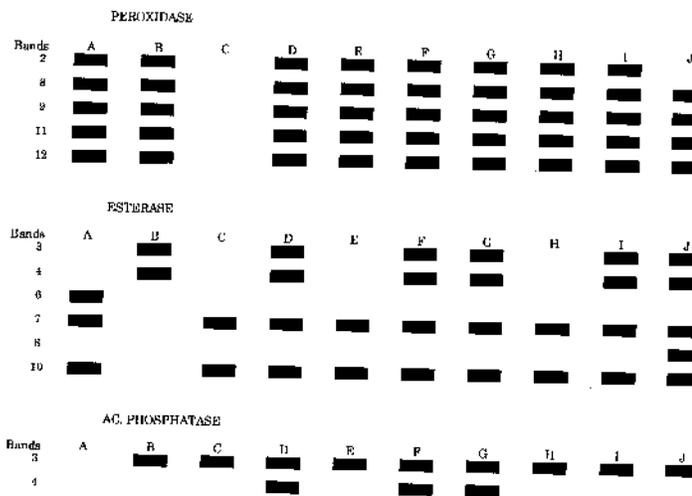


Fig. 1. Schematic zymograms of peroxidase, esterase, and acid phosphatase isozymes in female pistachio cultivars: A = 'Bianca'; B = 'Cappuccia'; C = 'Cerasola'; D = 'Ghiandalora'; E = 'Giulla'; F = 'Insolia'; G = 'Pignatone'; H = 'Silvana'; I = 'Tardiva'; J = 'Kerman'.

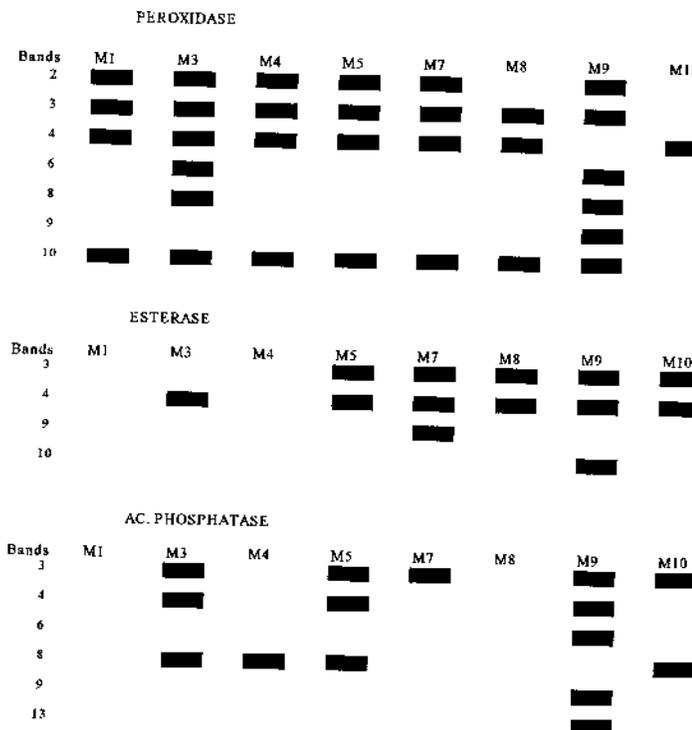


Fig. 2. Schematic zymograms of peroxidase, esterase, and acid phosphatase isozymes in male pistachio selections.

and 'Kerman') were uniquely identified with this enzyme system.

Peroxidases. Three female and five male groups of selections were evident. This system permitted full discrimination for 'Cerasola' and 'Kerman'. Among the males, differing patterns were shown by selections M3, M8, M9, and M10.

Acid phosphatases. AcPh distinguished six uniform groups of male selections and only three groups of female cultivars. With this system only 'Bianca' could be discerned among the females. Male selections M4, M7, M9, and M10 were uniquely identified.

Using these three enzyme systems together, it was possible to identify all eight male selections, but only these five of the 10 female cultivars: 'Bianca', 'Cappuccia', 'Cerasola', 'Kerman', and 'Tardiva', thus indicating a more pronounced degree of polymorphism among the male than among the female genotypes. Hence, a relative degree of similarity among the Sicilian pistillate cultivars is likely, as evidence from the dendrogram of the studied accessions, based on the results of cluster analysis, demonstrate (Fig. 3).

On the basis of these results, 'Cerasola' and 'Kerman' proved to be fairly distant from the other female cultivars, and M9 from the male selections. This result was expected, considering their extremely different origin and appearance. In contrast, a lower phylogenetic distance can be inferred between 'Gialla' and 'Silvana' and among 'Insolia', 'Pignatone', and 'Ghiandalora'.

Multivariate analysis. Canonical discriminant analysis for the female cultivars showed that a satisfactory degree (77%) of the total variance was summarized by the first three canonical functions; i.e., 44%, 23%, and 10%, respectively (Table 1). All CDFs resulted in significance at $P \leq 0.01$ by the likelihood test. As CDF_4 to CDF_9 accounted only for 23% of between-cultivar variation, its contribution was considered negligible; thus, only the first three CDFs were further examined. Fruit fresh weight, fruit dry weight, and fruit length had the greatest discriminatory power in the first two canonical functions, while apical leaflet width and leaf rachis length (RALE) loaded more on the third function. Nevertheless, the correlation coefficients obtained did not allow us to discern clearly the relative contribution of each character on CDFs (Table 1).

With these three functions, it was possible to clearly separate 'Kerman' and 'Silvana' from the others, since they are located at the opposite extremes of function 1 (Fig. 4A). Furthermore, ANOVA of canonical scores confirmed this clear discrimination (Table 2). 'Pignatone', 'Insolia', and 'Gialla' form a compact group closely surrounding 'Bianca'. Separated from this group are 'Cerasola', mainly for the second function; 'Tardiva' and 'Cappuccia', both for the first function and the third one; and, partially, 'Ghiandalora', for the third function.

About 93% of the total variance in the male selections was summarized with the first three canonical functions; i.e., 59%, 21%, and 13%, respectively (Table 3). All CDFs resulted in

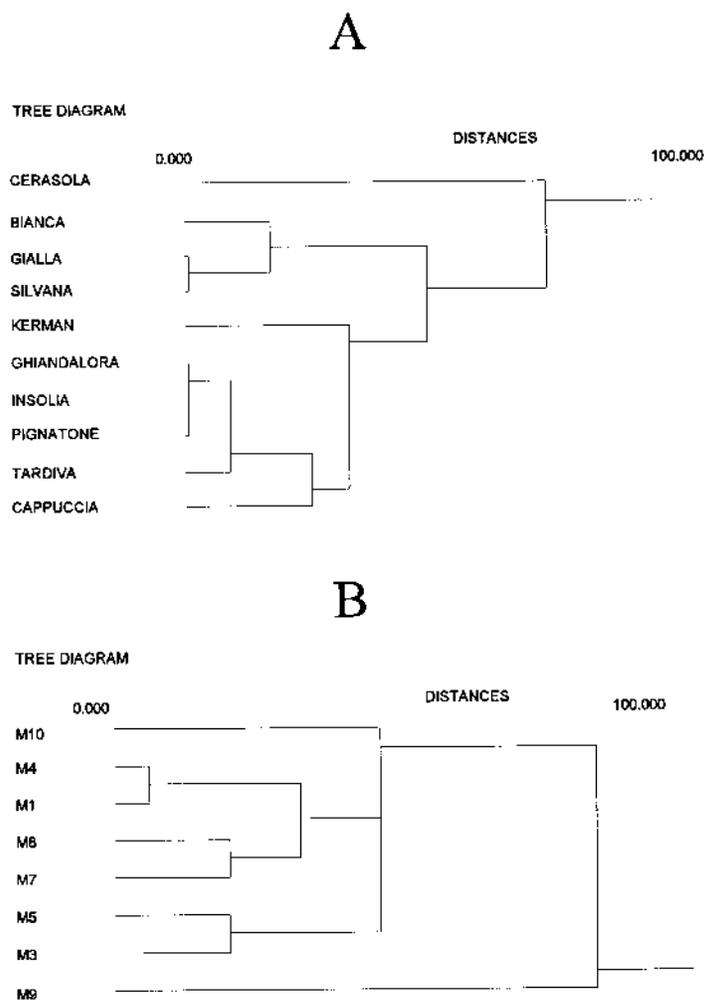


Fig. 3. Dendrogram of the studied accessions based on isozyme analyses. Metric distance calculated by cluster analysis is normalized percent disagreement (complete linkage method—farthest neighbor).

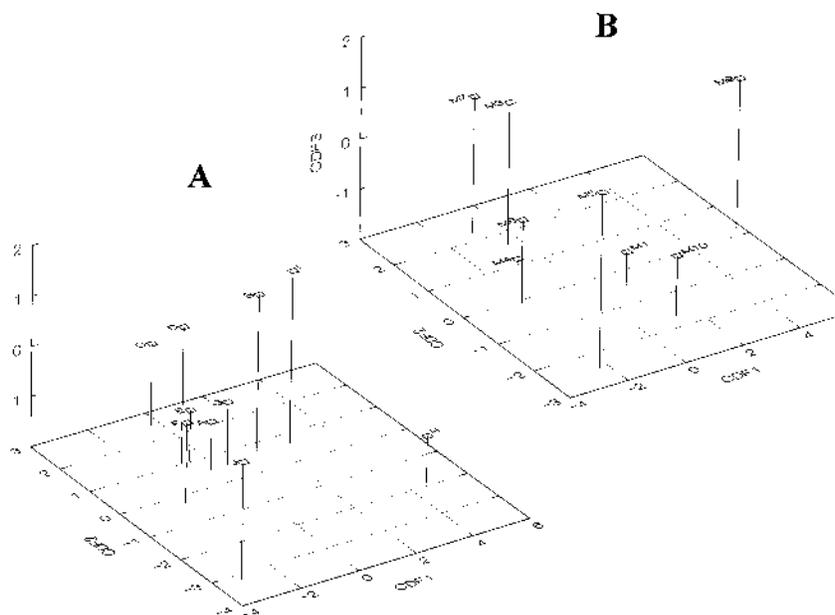


Fig. 4. (A) Means of standardized canonical scores of the first three canonical discriminant functions (CDF) for 10 female pistachio cultivars: A = 'Bianca'; B = 'Cappuccia'; C = 'Cerasola'; D = 'Ghiandalora'; E = 'Gialla'; F = 'Insolia'; G = 'Pignatone'; H = 'Silvana'; I = 'Tardiva'; J = 'Kerman'. (B) Means of standardized canonical scores of the first three CDFs for eight male pistachio selections.

significance at $P \leq 0.01$ by the likelihood test, except for CDF₇. The dimensions of the leaves, above all RALE, were the most separating characters. Particularly, from negative to positive values of function 1, RALE tended to increase; basal leaflet width and the dimensions of inflorescence bud loaded more on CDF₂; inflorescence bud fresh weight and central leaflet width loaded more on CDF₃. The male selection M9 was clearly separated by the plot of the three CDFs (Fig. 4B, Table 4), while the remaining selections present a scattered distribution, particularly along the axis corresponding to CDF₂. Thus, almost all of these selections may be considered to be fairly distinct from one another.

The discriminant power of the canonical functions obtained, as tested by checking predicted vs. actual group membership, was >70% in almost all the accessions and particularly high (>90%) for some female ('Kerman', 'Silvana', and 'Tardiva') and male (M5, M8, and M9) selections (Table 5). Such values can be considered satisfactory, since values of only 50% have been considered effective for discriminating between cultivars (Alessandri and Casini, 1992; Ueda and Okada, 1994). Our results are consistent with those of isozyme analysis for 'Kerman', 'Cerasola', and 'Tardiva', which produced distinct patterns. 'Ghiandalora', 'Insolia', and 'Pignatone' formed a compact group. Furthermore, CDA allowed us to discriminate between 'Gialla' and 'Silvana', which were not distinguished by isozymes.

These results demonstrate that combined use of CDA and isozyme analysis is useful for a rapid screening of *P. vera* germplasm and evaluation of variation among male and female germplasm. This isozyme analysis, based on the polymorphism of only three enzymes, allowed identification of 50% of the female cultivars and all male selections. A similar percentage of identified accessions was obtained with CDA, although with variation in the group of recognized cultivars. Nevertheless, the combination of the two methods enhanced the possibility of uniquely identifying the female cultivars. Identification was difficult or impossible for the compact group of 'Insolia', 'Pignatone', and 'Ghiandalora'. Nevertheless, Dollo (1991) reported a differing isozyme pattern of PGI for 'Insolia' and 'Ghiandalora'; thus, for 'Pignatone', possible synonymy with 'Insolia' or 'Ghiandalora' cannot be excluded. Together, CDA and the rare enzyme patterns in 'Kerman', 'Cerasola', and male selection M9 confirm the greater phylogenetic distance of these genotypes from the other accessions. 'Kerman', the primary American cultivar obtained from an Iranian seedling, belongs to a distinct geographical pool (Joley, 1969); 'Cerasola' is characterized by an extraordinary red pigmentation of the hull, similar to that of 'Red Aleppo'; while M9, locally known as 'Santangilisi', is a putative hybrid between *P. vera* and *P. terebinthus* (Bonifacio, 1942). The more pronounced degree of polymorphism for staminate than for pistillate accessions is probably due to a more intense selection pressure exerted by growers

Table 3. Standardized canonical coefficient (SCC) and correlation coefficient (r) between the first three canonical discriminant functions (CDF₁, CDF₂, CDF₃) and variables of morphometric characters of the male selections.

Character	Canonical discriminant function					
	CDF ₁		CDF ₂		CDF ₃	
	SCC	r	SCC	r	SCC	r
Leaf rachis length	0.800	0.486	-0.150	0.029	0.178	0.167
Basal leaflet length	0.080	0.017	0.069	0.111	-0.795	-0.073
Basal leaflet width	-0.439	-0.294	0.828	0.311	-0.044	0.286
Central leaflet length	0.387	0.067	-0.267	-0.117	0.187	0.331
Central leaflet width	-0.276	-0.292	0.110	0.105	0.836	0.542
Apical leaflet length	0.630	0.013	-0.371	-0.232	0.109	0.305
Apical leaflet width	-0.769	-0.271	-0.275	-0.113	-0.024	0.418
Inflorescence bud length	-0.346	0.066	-0.047	0.322	-0.979	-0.263
Inflorescence bud max diameter	0.087	0.302	0.331	0.579	-0.013	0.043
Inflorescence bud min diameter	0.184	0.256	0.476	0.586	-0.080	0.031
Inflorescence bud fresh weight	0.309	0.238	-0.062	0.443	0.991	0.134
Inflorescence bud dry weight	0.134	0.246	0.372	0.406	-0.038	0.046
Variance explained (%)	58.58		20.79		13.31	

Table 4. Means of standardized canonical scores of the first three canonical discriminant functions (CDF) for the male selections.

Male selection	Canonical discriminant function		
	CDF ₁ ^z	CDF ₂ ^z	CDF ₃ ^z
M1	1.017 c	-0.513 c	-1.283 ab
M3	-0.941 b	1.255 d	1.094 de
M4	-0.878 b	1.013 d	-1.895 a
M5	-2.311 a	-2.450 a	1.459 e
M7	-0.986 b	2.218 e	0.726 d
M8	-2.356 a	-0.300 c	-0.189 c
M9	5.804 d	0.190 c	0.991 de
M10	1.319 c	-1.677 b	-0.805 bc

^zMean separation within each column by Tukey's HSD test ($P \leq 0.01$).

on the female trees and to the lack of seed propagation that would confirm the statements of many authors that vegetative propagation is the only propagation method used by growers (Bonifacio, 1942; Minà Palumbo, 1882). The existence of a rare isozyme pattern, if substantiated by further research, might suggest the existence of two distinct genetic pools within the pistachio germplasm of Sicily. This hypothesis could be explained by sequential introductions of genetic material, the reddish 'Cerasola' from Syria by the Romans and the later introductions by the Arabs. This sequence would confirm Maggs' (1973) statement that Sicily can be regarded as a late, secondary, center of pistachio genetic diversity.

Further investigation, extended to other genotypes from various countries and other enzyme systems or DNA analysis techniques, not susceptible to environmental influences, would answer this question.

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Table 5. Percentage of correctly predicted group membership of female and male accessions back-classified using the first three canonical discriminant functions.

Accession	Cases correctly classified (%)
Female cultivar ^z	
A	74
B	79.6
C	76
D	74
E	70
F	85.8
G	63.3
H	94
I	96
J	100
Male selection	
M1	80.9
M3	70
M4	81.3
M5	91.1
M7	69.4
M8	90
M9	97.7
M10	76.6

^zA = 'Bianca'; B = 'Cappuccia'; C = 'Cerasola'; D = 'Ghiandalora'; E = 'Gialla'; F = 'Insolia'; G = 'Pignatone'; H = 'Silvana'; I = 'Tardiva'; J = 'Kerman'.

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